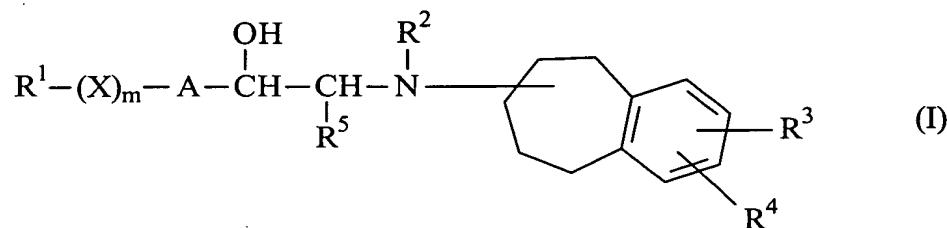


IN THE CLAIMS

Please add Claims 36-48:

--36 (New) A compound of the general formula (I):



wherein

C

R¹ is aryl which may have one or more suitable substituent(s), heterocyclic group or cyclo(lower)alkyl,

R² is hydrogen or amino protective group,

R³ and R⁴ are independently hydrogen, halogen, hydroxy, amino, nitro, carboxy, protected carboxy, aryl, lower alkyl, hydroxy(lower)alkyl, amino(lower)alkyl, acyloxy(lower)alkyl, acylamino(lower)alkyl, lower alkylamino(lower)alkyl which may have one or more suitable substituent(s), mono or di-(lower)alkylamino, acylamino, acyl group, lower alkoxy, halo(lower)alcoxy, lower alkenyloxy, lower alkoxy(lower)alcoxy, aryloxy, cyclo(lower)alkyloxy, heterocyclicoxy, ar(lower)alkyloxy, acyloxy, lower alkylcarbamoyl(lower)alcoxy, heterocycliccarbamoyl(lower)alcoxy, heterocycliccarbonyl(lower)alcoxy, N-lower alkyl-lower alkylcarbamoyl(lower)alcoxy, arylcarbamoyl(lower)alcoxy which may have lower alkoxy or di(lower)alkylamino, di-lower alkylsulfonyloxy, N-lower alkyl-heterocyclic(lower)alkylcarbamoyl(lower) alcoxy, N-lower alkyl-lower alkylcarbamoyl(lower)alcoxy or N-lower alkyl-cyclo(lower)alkylcarbamoyl(lower)alcoxy,

R⁵ is hydrogen, lower alkyl, or aryl,

A is lower alkylene which may have one or more suitable substituent(s) or lower alkenylene,

X is O, S, SO, SO₂ or NH, and

m is an integer of 0 or 1,

or a salt thereof,

wherein when R¹ is naphthyl and R⁵ is H, then X is not O.

24 37. (New) The compound of claim 36, wherein

R¹ is phenyl which may have 1 or 2 suitable substituent(s) selected from the group consisting of hydroxy and lower alkylsulfonylamino,

R² is hydrogen,

R³ is lower alkylcarbamoyl(lower)alkoxy, heterocycliccarbamoyl(lower)alkoxy, heterocycliccarbonyl(lower)alkoxy, N-lower alkyl-lower alkylcarbamoyl(lower)alkoxy, hydroxy, lower alkoxy, protected carboxy, arylcarbamoyl(lower)alkoxy which may have lower alkoxy or di(lower)alkylamino, di-lower alkylsulfamoyloxy, N-lower alkyl-heterocyclic(lower)alkylcarbamoyl(lower) alkoxy, N-lower alkyl-lower alkylcarbamoyl(lower)alkoxy or N-lower alkyl-cyclo(lower)alkylcarbamoyl(lower)alkoxy,

R⁴ is hydrogen,

R⁵ is hydrogen,

A is lower alkylene,

X is O, and

m is an integer of 1--

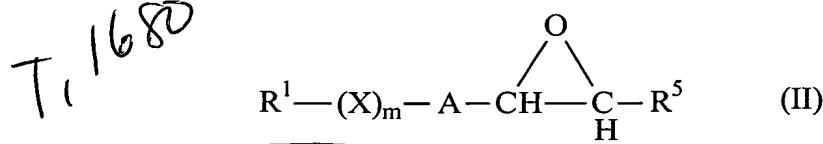
24 38. (New) The compound of claim 37, wherein

R¹ is phenyl which may have hydroxy and methylsulfonylamino,

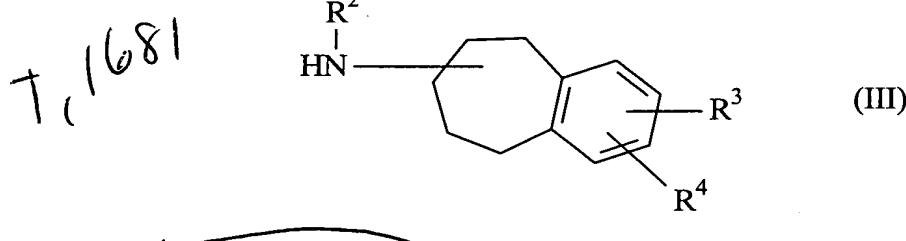
R^3 is ethylcarbamoylmethoxy, indolylcarbamoylmethoxy, piperidinocarbonylmethoxy, N-methylbutylcarbamoylmethoxy, hydroxy, butylcarbamoylmethoxy, methoxy, methoxycarbonyl, ethoxy, dimethylsulfamoyloxy, tetrazolylcarbamoylmethoxy, N-methylpyridylethylcarbamoylmethoxy, methoxyphenylcarbamoylmethoxy, thiazolylcarbamoylmethoxy, dihydroindolylcarbamoylmethoxy, N-ethylpropylcarbamoylmethoxy, N-methylbutylcarbamoylmethoxy, N-ethylbutylcarbamoylmethoxy, dimethylaminophenylcarbamoylmethoxy or N-methylcyclohexylcarbamoylmethoxy.

39. (New) A process for preparing a compound of claim 36, or a salt thereof, which comprises,

(i) reacting a compound (II) of the formula:

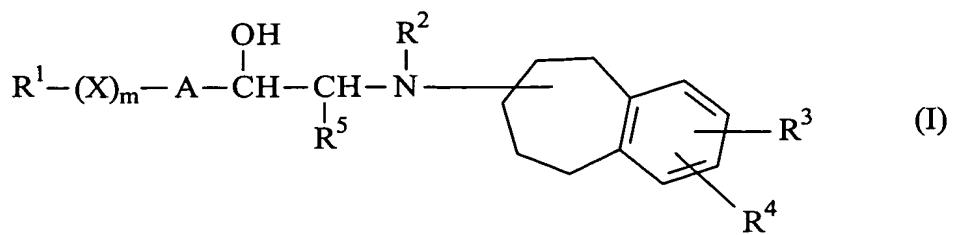


wherein R^1 , R^5 , A , X and m are each as defined in claim 36, with a compound (III) of the formula:



wherein R^2 , R^3 and R^4 are each as defined in claim 36, or a salt thereof, to give a compound (I) of the formula:

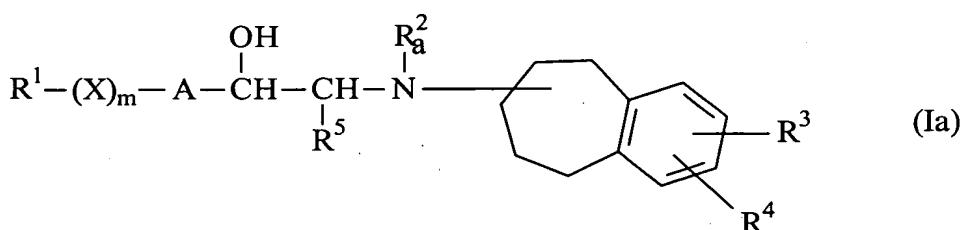
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wherein R^1 , R^2 , R^3 , R^4 , R^5 , A, X and m are each as defined in claim 36, or a salt thereof, or

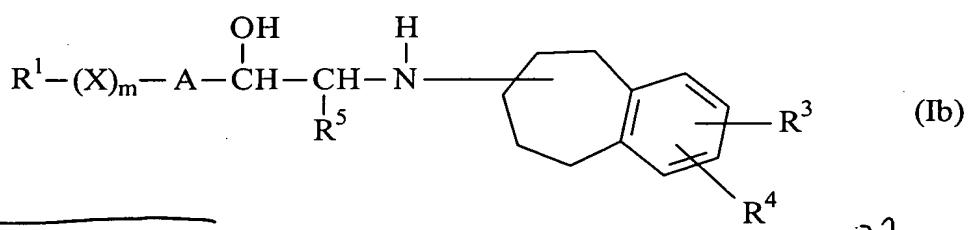
(ii) subjecting a compound (Ia) of the formula:

Ti¹⁶⁹
Cr



wherein R^1 , R^3 , R^4 , R^5 , A, X and m are each as defined in claim 36, and R_a^2 is amino protective group, or a salt thereof, to elimination reaction of the amino protective group, to give a compound (Ib) of the formula:

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wherein R¹, R³, R⁴, R⁵, A, X and m are each as defined in claim 36, or a salt thereof.

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4Q. (New) A pharmaceutical composition which comprises the compound of claim 36 or a pharmaceutically acceptable salt thereof in admixture with a pharmaceutically acceptable carrier or excipient.

32
41. (New) A method for making a pharmaceutical composition or a medicament comprising admixing the compound of claim 36 or a pharmaceutically acceptable salt thereof with a pharmaceutically acceptable carrier or excipient.

33
42. (New) A compound of claim 36 or a pharmaceutically acceptable salt thereof in the form of a tablet, pellet, troche, capsule, suppository, cream, ointment, aerosol, powder for insufflation, solution, emulsion, or suspension.

34
43. (New) A method for the prophylactic and/or the therapeutic treatment of pollakiuria or urinary incontinence which comprises administering an effective amount of a compound of claim 36 or a pharmaceutically acceptable salt thereof to a subject in need thereof.--

35
44. (New) A method for agonizing a β_3 adrenergic receptor comprising contacting said receptor with the compound of claim 36.

36
45. (New) A method for inducing gut-selective sympathomimetic activity comprising administering an effective amount of the compound of claim 36 to a subject in need thereof.

37
46. (New) A method for the prophylactic and/or the therapeutic treatment of a gastrointestinal disorder comprising administering an effective amount of the compound of claim 36 or a pharmaceutically acceptable salt thereof to a subject in need thereof.

38
47. (New) A method for the prophylactic and/or the therapeutic treatment of an ulcer or pancreatitis comprising administering an effective amount of the compound of claim 36 or a pharmaceutically acceptable salt thereof to a subject in need thereof.

39
48. (New) A method for inducing lysis comprising administering an effective amount of the compound of claim 36 or a pharmaceutically acceptable salt thereof to a subject in need thereof.----